



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s): Ellen W. Evans

Application No.: 10/673,888

Filed: 09/29/2003

For: Methods for Treating Disorders of Calcium Homeostasis

Group Art Unit: 1623

Examiner: Roy P. Issac

Atty. Docket No. : OC01600 US

X

INFORMATION DISCLOSURE STATEMENT TRANSMITTAL

Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450
Mail Stop: Amendment
Sir:

(a) Pursuant to the duty of disclosure under 37 C.F.R. § 1.56, it is requested that the documents listed on the accompanying Forms PTO/SB/08a & 08b be considered and made of record in the above-identified patent application. Other than cited U.S. patents and published patent applications, copy(ies) of these documents ☒ are enclosed ☐ were filed in related U.S. Application No(s) _____ filed _____ respectively. Since the Examiner is believed to have ready access to the cited U.S. patents and published patent applications, copies of these documents are not herewith transmitted, but will be provided upon request.

(b) No fee is believed to be due because:

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- ☐ This Information Disclosure Statement is being submitted concurrently with a Request for Continued Examination (RCE); **or**
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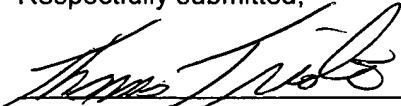
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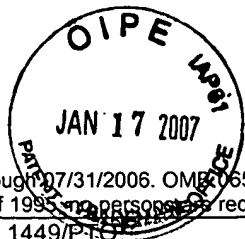
Dated: 01/17/2007

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Respectfully submitted,

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PTO/SB/08B (08-03)

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (USE AS MANY SHEETS AS NECESSARY)		ATTY. DOCKET NO.: OC01600 US		SERIAL NO.: 10/673,888	
		APPLICANT: Evans <i>et al.</i>			
		FILING DATE: Sept. 29, 2003		GROUP: 1623	
		OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)			
Examiner initials*	Cite no. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.			T ²
	AA	FISHER <i>et al.</i> , Alendronate mechanism of action: geranylgeraniol, an intermediate in the mevalonate pathway, prevents inhibition of osteoclast formation, bone resorption, and kinase activation in vitro, Proc Natl Acad Sci U S A. 1999 Jan 5;96(1):133-8.			
	AB	DUNFORD <i>et al.</i> , Structure-activity relationships for inhibition of farnesyl diphosphate synthase in vitro and inhibition of bone resorption in vivo by nitrogen-containing bisphosphonates, J Pharmacol Exp Ther. 2001 Feb;296(2):235-42.			
	AC	OLGAARD <i>et al.</i> , Prevention of uremic bone disease using calcimimetic compounds, Annu Rev Med. 2001;52:203-20.			
	AD	FOX <i>et al.</i> , Calcimimetic compound NPS R-568 stimulates calcitonin secretion but selectively targets parathyroid gland Ca(2+) receptor in rats, J Pharmacol Exp Ther. 1999 Aug;290(2):480-6.			
	AE	CHATTOPADHYAY <i>et al.</i> , Cellular "sensing" of extracellular calcium (Ca(2+)(o)): emerging roles in regulating diverse physiological functions, : Cell Signal. 2000 Jun;12(6):361-6.			
	AF	COBURN <i>et al.</i> , Calcium-sensing receptor and calcimimetic agents. Kidney Int Suppl. 1999 Dec;73:S52-8.			
	AG	CANALIS, Novel treatments for osteoporosis, J Clin Invest. 2000 Jul;106(2):177-9.			
	AH	GOODMAN <i>et al.</i> , The Calcimimetic agent AMG 073 lowers plasma parathyroid hormone levels in hemodialysis patients with secondary hyperparathyroidism, J Am Soc Nephrol. 2002 Apr;13(4):1017-24.			
	AI	NEMETH <i>et al.</i> , Calcimimetics with potent and selective activity on the parathyroid calcium receptor, Proc Natl Acad Sci U S A. 1998 Mar 31;95(7):4040-5.			
	AJ	COHEN <i>et al.</i> , Inhibitors of prenylation of Ras and other G-proteins and their application as therapeutics, Biochem Pharmacol. 2000 Oct 15;60(8):1061-8.			
Examiner signature:		Date considered:			

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